PATENT COOPERATION TREATY

PCT

TRANSLATION INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 443520EH			nce	FOR FURTHER A	ACTION	See Form PCT/IPEA/416		
Internati	onal app	olication No.		International filing da	ate (day/month/year)	Priority date (day/month/year)		
PCT/EP2004/012988			2988	16.11.200	4	20.11.2003		
Internati	onal Pat	ent Classification	on (IPC) or nati	onal classification and	IPC			
G01	G01N33/543							
Applicant NOVEMBER AKTIENGESELLSCHAFT								
1.	1. This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.							
2.	This R	EPORT consist	s of a total of	8	sheets, including this cover sheet.			
3.	This re	port is also acc	ompanied by A	NNEXES, comprising:	:			
	a	sent to the	e applicant and	to the International Bi	ureau) a total of	sheets, as follows:		
	sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).							
	sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental							
	, _	Box.	T	D 1) 4 1 6		6.1.4		
	b. <u></u>	(sent to the	e International i	Bureau oniy) a totai oi	(indicate type and number	r of electronic carrier(s))		
	, containing a sequence listing and/or tables related thereto, in computer readable form only, as indicated in the Supplemental Box Relating to Sequence Listing (see							
				rative Instructions).	is indicated in the Supplei	mental Box Relating to Sequence Listing (see		
4.	This re	port contains in	dications relati	ng to the following iter	ns:			
	\boxtimes	Box No. I	Basis of the	report				
		Box No. II	Priority					
		Box No. III	Non-establi	shment of opinion with regard to novelty, inventive step and industrial applicability				
	Box No. IV Lack of unit		y of invention					
	Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement							
		Box No. VI	Certain doc	uments cited				
		Box No. VII	Certain defe	ects in the international	application			
	Box No. VIII Certain observations on the international application							
Date of submission of the demand				Date of completion of thi	is report			
Name and mailing address of the IPEA/EP				Authorized officer				
Facsimile No.				Telephone No.				

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International application No.

PCT/EP2004/012988

Box	No. I	Basis of the report								
1.		n regard to the language, this report is based on the internationated under this item.	onal application in the language in which	it was filed, unless otherwise						
		This report is based on translations from the original language which is the language of a translation furnished for the purp		,						
		international search (Rule 12.3 and 23.1(b))								
		publication of the international application (Rule 12.4	4)							
		international preliminary examination (Rule 55.2 and	,							
2.	recei		rd to the elements of the international application, this report is based on (<i>replacement sheets which have been furnished to the Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to):</i>							
		the international application as originally filed/furnished								
	\boxtimes	the description:								
		pages 1-17		as originally filed/furnished						
		pages*	received by this Authority on							
		pages*	received by this Authority on							
	\boxtimes	the claims:								
		nos. 1-30		as originally filed/furnished						
		nos.*	as amended (together with	any statement) under Article 19						
		nos.*	received by this Authority on							
		nos.*	received by this Authority on							
	\boxtimes	the drawings:								
		sheets 1/5-5/5		as originally filed/furnished						
		sheets*	received by this Authority on							
		sheets*	received by this Authority on							
		a sequence listing and/or any related table(s) – see Supplem	nental Box Relating to Sequence Listing.							
3.		The amendments have resulted in the cancellation of:								
		the description, pages								
		the claims, nos.								
		the decivings shoots/figs								
		the sequence listing (specify):								
		any table(s) related to sequence listing (specify):								
4.		This report has been established as if (some of) the amend they have been considered to go beyond the disclosure as fi								
		the description, pages								
		the claims, nos.								
		the drawings, sheets/figs	the drawings, sheets/figs							
		the sequence listing (specify):								
		any table(s) related to sequence listing (specify):								
*	If ite	m 4 applies, some or all of those sheets may be marked "sup	erseded."							

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Box No. V		Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1.	Statement					
	Novelty	(N) Claims 1-12, 14-30	YES			
		Claims 13	_ NO			
	Inventiv	ve step (IS) Claims 1-12	YES			
		Claims 13-30	_ NO			
	Industri	al applicability (IA) Claims 1-30	YES			
		Claims	_ NO			
2.	Citations	nd explanations (Rule 70.7)				
۷.	1.	This report makes reference to the following				
	⊥•	documents:				
		documents.				
	D1:	WO 02/43865 A				
	D2:	WO 03/049530 A				
	D3:	DE 101 11 520 A1				
	D4:	US 2003/095897 A1				
	D5:	WO 2004/096443 A (NOVEMBER AG GESELLSCHAFT FÜR				
		MOLEKULARE MEDIZIN; BART), 11 November 2004 (2004-				
		11-11)				
	2.	Novelty (PCT Article 33(1) and 33(2))				
	2.1	Claim 13:				
		Document D1, which is considered to represent the				
		closest prior art, discloses a device which				
		comprises:				
		• a first container (figure 3, [3]) for providing				
		or receiving a first liquid and paramagnetic				
		microparticles (abstract);				
		• a first pipe that opens into the first container				
		(figure 1, [15, 13]); and				
		 a section of the first pipe which, in relation 				

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to the remaining cross-sectional area of the first pipe, has an enlarged cross-sectional area (figure 1, [15, 13]);

- a first magnet or a first recess for receiving a first magnet for generating a first magnetic field in a zone of the first container and in the pipe section (figure 3); or
- a first magnet or a first recess for receiving a first magnet for generating a first magnetic field in a zone of the first container, and a second recess for receiving a second magnet for generating a second magnetic field in the pipe section (figure 3);
- the zone, the pipe section and the first recess or first magnet, as well as, if provided, the second recess, are arranged and/or shaped in such a way that the magnetic field inside the pipe section has a stronger average field intensity than the magnetic field inside the first container (figure 3).

The subject matter of claim 13 is therefore not novel. It should also be noted that, owing to the use of "or" in claim 13, the second magnet is not a feature of the device claim.

It should also be noted that the difference from the device disclosed in D4 (claim 55), in comparison with claim 13, is that no enlarged section in relation to the section of the first pipe is disclosed. However, D4 discloses the

PCT/EP2004/012988 Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement arrangement of a plurality of magnets along the liquid path. 3. Inventive step (PCT Article 33(1) and 33(3)) 3.1 Claim 1: Document D1, which is considered to represent the closest prior art, discloses improved purification processes for a first substance (abstract) bound to paramagnetic microparticles (abstract), the microparticles being suspended in a first liquid (claim 1). The processes have the following steps: (a) the microparticles are exposed in a first container to a first magnetic field in order to retain them and prevent them from being washed away with a stream of a first liquid (claim 1, figure 1); and (b) at least part of the first liquid is led after step (a) in a first direction through a first pipe, through a section (figure 1, [5]) of the first pipe. The subject matter of claim 1 differs from the above in that

> part of the first liquid in the section (figure 1, [5]) is exposed to a second or to the first magnetic field in order to retain microparticles which have nevertheless been washed away, the cross-sectional area of the first pipe being enlarged in the section, and the second or first

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magnetic field inside said section having a

magnetic field inside said section having a stronger average field intensity than the first magnetic field inside the first container.

The subject matter of claim 1 is thus novel (PCT Article 33(2)).

The present invention can therefore be considered to address the problem of devising a particularly effective process for retaining paramagnetic particles.

The solution to this problem, as proposed in claim 1 of the present application, involves an inventive step (PCT Article 33(3)) for the following reasons:

D1 proposes a process for conveying and concentrating magnetic particles. A solution A, in which the analyte is bound to magnetic particles, is poured into a vessel (1) linked to a second vessel by a pipe. The analyte bound to the magnetic particles is conveyed into the second vessel by means of a magnet. D1 does not use the first magnet or a second magnet at another location in the delivery pipe or in a second vessel in order to retain the microparticles.

D2 describes a process for washing magnetic particles in which the magnetic particles are led through a throughflow cell and retained by a magnet in a first zone, then the magnetic field is

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switched off and the magnetic particles are stirred. Stirring can be carried out by switching the magnetic field on and off. D2 does not use the first magnet or a second magnet at another location of a delivery pipe or in a second vessel in order to retain the microparticles.

D3 describes a process and device for purifying biomolecules by means of magnetic particles. Firstly, magnetic particles are added to the starting solution, and the starting solution is mixed. The magnetic particles are prepared in such a way that they selectively bond to biomolecules in the resultant suspension, either immediately or after admixture of a binding buffer. The suspension is then introduced into a flexible tube that goes through a strong magnetic field. The particles are retained therein and can be washed. After the magnetic field is removed, the particles can be suspended again. That process is characterised by handling in a metering system comprising canulae, flexible tube and metering syringe (dilutor), the suspension being led in a flexible tube through the magnetic field. The magnetic particles are purified with various washing buffers in the flexible tube of the metering system, permitting even the smallest volumes to be handled without losses. The embodiment described permits multiple washing buffers to be received in the flexible tube before the suspension, and the particles to be washed without having to go through separate vessels

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containing washing solutions.

D4 describes a process in which a mixture of magnetic particles is led past a magnet in one direction, then led during a subsequent step in another direction past the magnet (claim 37). D4 also describes an apparatus having a plurality of magnetic fields arranged at a fixed location of a liquid path (claim 55). D4 does not describe different cross-sectional areas in an additional pipe, besides the first pipe, or a second or first magnetic field having a stronger average field strength inside the pipe section.

Either alone or in combination, the documents fail to suggest the claimed process. The subject matter of claim 1 and of its dependent claims is thus inventive in relation to the citations.

3.2 Claims 14-30:

Dependent claims 14-30 do not contain any features which, in combination with the features of any claim to which they refer, meet the PCT inventive step requirements.